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(54) MÉTHODE DE PRÉPARATION DE POLYMERIS D'ETHYLENE
(54) PROCESS FOR THE PREPARATION OF ETHYLENE
POLYMERS

(52) The preparation of polyethylene and ethylene-1-olefin copolymers in the suspension or gas phase process with the aid of metalalkoxo/alkylene oxide catalysts where metallocene component is a bridged bis(cyclopentadienyl) complex offers particular advantages. Depending on the structure of the complex, the use of these complexes gives, with high activity, products in a very broad molecular-weight range and permits the production of very varied grain morphologies of the products, such as high and low bulk density, extremely small and extremely large mean grain diameter and various grain shapes. The variety of grain morphologies which can be achieved opens up various possibilities for the use of such polyethylene products in molding processes.



Industry Canada Industry Canada



MOLCUST ANTILLES/ULLSCHAND

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Description

Process for the preparation of ethylene polymers

5 The present invention relates to a process for the preparation of polyethylene and ethylene-1-olefin copolymers of various molecular weight ranges with the aid of metallocene/aluminoxane catalysts.

10 A process for the preparation of polyethylenes with the aid of metallocene/aluminoxane catalysts in toluene as suspending agent has already been described (cf. EP 69,951). The molecular weights achieved for the polymers are relatively low. No data is given on the morphology of the polymers.

15 Furthermore, a comparable process using a high-boiling hydrocarbon as suspending agent has been disclosed (cf. EP 170,059). However, the catalyst activities are moderate, and the bulk densities achieved for the polymer are between 0.15 and 0.18 g/cm³.

20 Furthermore, processes have been described for the preparation of polyethylene and ethylene-1-olefin copolymers with the aid of metallocene/aluminoxane catalysts by polymerization in the gas phase (cf. EP 206,794, 205,443 and 204,842). Here too, only moderately high molecular weights of the polymers and in most cases only poor 25 activities of the catalysts are achieved.

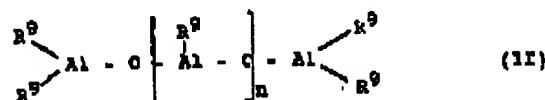
25 It is common to all the abovementioned processes that, as the metallocene, unbridged bicyclopentadienylmiconium complexes are employed in which the cyclopentadienyl radicals are substituted or unsubstituted, and the metallocene is either employed as such in the polymerization or has been linked to an inert support by appropriate preliminary reaction steps.

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It has now been found that the preparation of polyethylene and ethylene-1-olefin copolymers by suspension or gas-phase processes with the aid of metallocene/aluminoxane catalysts whose metallocene component is a bridged bis(cyclopentadienyl) complex offers important advantages.

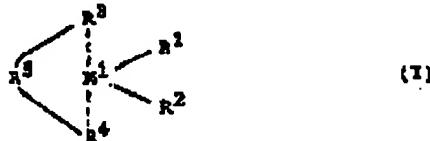
The invention thus relates to a process for the preparation of ethylene polymers by polymerisation of ethylene or copolymerisation of ethylene with 1-olefins having 3 to 20 carbon atoms at a temperature of from -60 to 200°C and a pressure of from 0.5 to 200 bar in solution, in suspension or in the gas phase, in the presence of a catalyst comprising a metallocene as the transition-metal component and an aluminoxane of the formula II



for the linear type and/or of the formula III



for the cyclic type, where, in the formulas II and III, R⁹ is a C₂-C₄-alkyl group or phenyl or benzyl, and n is an integer from 2 to 50, which comprises carrying out the polymerisation in the presence of a catalyst whose transition-metal component is a compound of the formula I



in which

M¹ is titanium, zirconium, hafnium, vanadium, niobium or tantalum,
R¹ and R² are identical or different and are a hydrogen

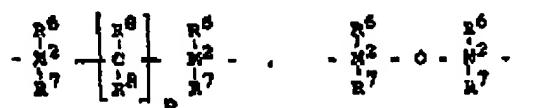
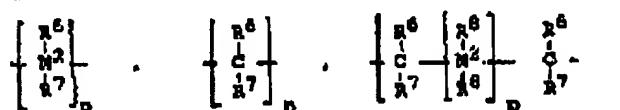
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atom, a halogen atom, a C_1 - C_{18} -alkyl group, a C_1 - C_{18} -alkoxy group, a C_1 - C_{18} -aryl group, a C_6 - C_{18} -aryloxy group, a C_6 - C_{18} -alkenyl group, a C_6 - C_{18} -arylalkyl group, a C_6 - C_{18} -alkylaryl group or a C_6 - C_{18} -arylalkenyl group.

5 R^3 and R^4 are identical or different and are a mono-nuclear or polynuclear hydrocarbon radical which, with the central atom M^2 , is able to form a sandwich structure,

10 M^3 is



15 $-BR^6$, $-AlR^3$, $-Ge-$, $-Sn-$, $-O-$, $-S-$, $-SO$, $-SO_2$, $-MR^3$, $-CO$, $-PR^3$ or $-P(O)R^3$, where R^6 , R^3 and R^2 are identical or different and are a hydrogen atom, a halogen atom, a C_1 - C_{18} -alkyl group, a C_1 - C_{18} -fluoroalkyl group,

20 a C_6 - C_{18} -fluoroaryl group, a C_1 - C_{18} -aryl group, a C_1 - C_{18} -alkoxy group, a C_1 - C_{18} -alkenyl group, a C_6 - C_{18} -arylalkyl group, a C_6 - C_{18} -alkylaryl group or a C_6 - C_{18} -alkylalkenyl group, or R^6 and R^7 or R^3 and R^2 , in each case with the atoms connecting them, form a ring,

M^2 is silicon, germanium or tin, and

p is the number 1, 2, 3, 4 or 5.

25 The catalyst to be used for the process according to the invention comprises an aluminosilane and a metallocene of the formula I

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- R^6 , - R^6R^7 , - Ge^6 , - Ge^6- , - $\text{C}-$, - $\text{C}-$, - SO_2 , - SO_2R^6 , - NH^6 ,
 - CO , - PR^6 or - $\text{P}(\text{O})\text{R}^6$, where R^6 , R^7 and R^8 are identi-
 5 cal or different and are a hydrogen atom, a halogen
 atom, preferably chlorine, a $\text{C}_1\text{-C}_{10}$ -, preferably $\text{C}_1\text{-C}_4$ -
 alkyl group, in particular a methyl group, a $\text{C}_1\text{-C}_{10}$ -fluoroalkyl group, preferably a $\text{C}_1\text{-C}_4$ group, a $\text{C}_1\text{-C}_{10}$ -fluorophenyl
 10 group, preferably a pentafluorophenyl group, a $\text{C}_1\text{-C}_{10}$ -, preferably $\text{C}_1\text{-C}_4$ -aryl group, a $\text{C}_1\text{-C}_{10}$ -, preferably $\text{C}_1\text{-C}_4$ -alkoxy group, in particular a
 15 methoxy group, a $\text{C}_1\text{-C}_{10}$ -, preferably $\text{C}_1\text{-C}_4$ -alkenyl group, a $\text{C}_1\text{-C}_{10}$ -, preferably $\text{C}_1\text{-C}_4$ -alkylalkenyl group or a $\text{C}_1\text{-C}_{10}$ -, preferably $\text{C}_1\text{-C}_4$ -alkylaryl group, or R^6 and R^7
 20 or R^6 and R^8 in each case, together with the
 25 atoms connecting them, form a ring,
 M^2 is silicon, germanium or tin, preferably silicon or
 30 germanium, and
 p is the number 1, 2, 3, 4 or 5.

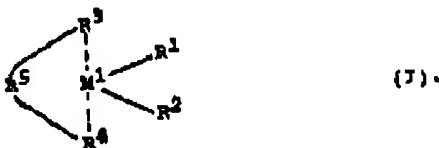
Particularly preferred metallocenes are:

20 rac-dimethylsilyl bis(1-indenyl)zirconium dichloride,
 rac-diphenylsilyl bis(1-indenyl)zirconium dichloride,
 1-milincyclobetyl bis(1'-indenyl)zirconium dichloride,
 rac-dimethylsilyl bis(1-(3-methylindenyl))zirconium
 25 dichloride,
 rac-1,1,2,2-tetramethylsilyl bis(1'-indenyl)zir-
 conium dichloride,
 dimethylsilyl bis(1-(3-trimethylsilyl)cyclopentadienyl)-
 zirconium dichloride, and
 30 diphenylmethylene(5-fluorophenyl)cyclopentadienylzirconium
 dichloride.

The above-described metallocenes can be prepared by the general reaction scheme below:

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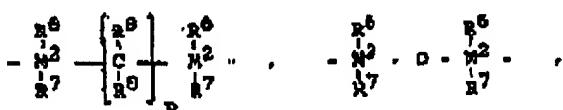
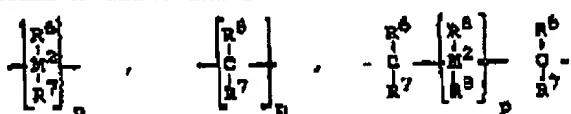
In the formula I, M¹ is a metal selected from the group comprising titanium, zirconium, hafnium, vanadium, niobium and tantalum, preferably zirconium.

5 R⁴ and R³ are identical or different and are a hydrogen atom, a C₁-C₁₀-, preferably C₁-C₅-alkyl group, a C₁-C₁₀-, preferably C₁-C₅-alkoxy group, a C₆-C₁₀-, preferably C₆-C₁₀-aryl group, a C₆-C₁₀-, preferably C₆-C₁₀-acryloxy group, a C₆-C₁₀-, preferably C₆-C₁₀-alkenyl group, a C₆-C₁₀-, preferably C₆-C₁₀-alkylalkyl group, a C₆-C₁₀-, preferably C₆-C₁₀-alkylaryl group or a halogen atom, preferably chlorine.

10 R³ and R⁴ are identical or different and are a mononuclear or polynuclear hydrocarbon radical which, together with the central atom M¹, is able to form a sandwich structure.

15 R³ and R⁴ are preferably cyclopentadienyl, indenyl, tetrahydroindenyl or fluorenyl, it also being possible for the basic structures to carry additional substituents.

20 R¹ is a mono- or polynumbered bridge which links the radicals R³ and R⁴ and is

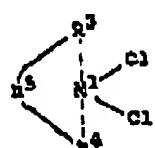


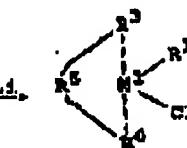
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$$\xrightarrow{\text{a butylli}}$$

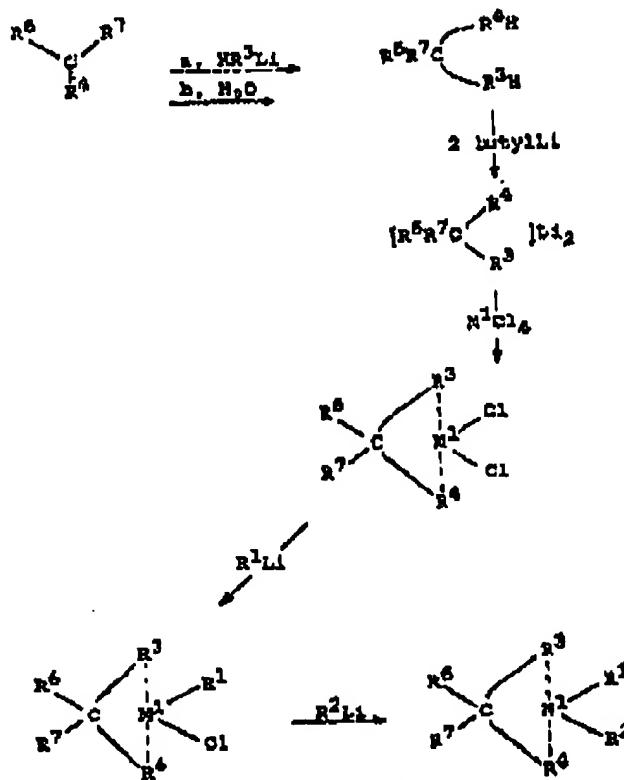

$$\xrightarrow{N^3LiA}$$


$$(X = Cl, Br, I, O-tosyl)$$


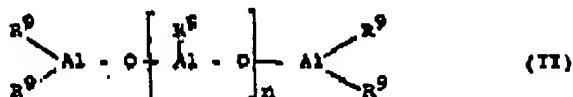
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xx



The cocatalyst is an aluminumane of the formula II



or the linear type and/or of the formula III

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for the cyclized type. In these formulas, R^9 is a C_1 - C_{20} alkyl group, preferably methyl, ethyl or isobutyl, butyl or neopentyl, or phenyl or benzyl. Methyl is particularly preferred. n is an integer from 2 to 50, preferably 3 to 10. However, the exact structure of the aluminoxane is not known.

The aluminoxane can be prepared in various ways.

One possibility is the careful addition of water to a dilute solution of a trialkylaluminum by introducing the solution of the trialkylaluminum, preferably trimethylaluminum, and the water in each case in small portions into an initially introduced larger amount of an inert solvent and between each portion awaiting the end of the evolution of gas.

In another process, finely powdered copper sulfate pentahydrate is slurried in toluene, and, in a glass flask under an inert gas at about -20°C , sufficient trialkylaluminum is added so that about 1 mole of $\text{CuSO}_4 \cdot 5\text{H}_2\text{O}$ is available for each 4 Al atoms. After slow hydrolysis with elimination of alkane, the reaction mixture is left at room temperature for 24 to 48 hours, cooling possibly being needed to ensure that the temperature does not exceed 30°C . The aluminoxane dissolved in the toluene is subsequently filtered off from the copper sulfate, and the solution is evaporated in *vacuo*. It is assumed that the low-molecular-weight aluminoxanes condense in the preparation process to form higher oligomers with elimination of trialkylaluminum.

Furthermore, aluminoxanes are obtained if trialkylaluminum, preferably trimethylaluminum, dissolved in an inert aliphatic or aromatic solvent, preferably heptane or

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toluene, is reacted at a temperature of from -20 to 100°C with aluminum salts, preferably aluminum sulfate, containing water of crystallization. In this case, the volume ratio between the solvent and the alkylaluminum 5 used is 1:1 to 30:1 - preferably 5:1 - and the reaction time, which can be monitored from the elimination of the alkyne, is 1 to 200 hours - preferably 10 to 40 hours.

10 Of the aluminum salts containing water of crystallization, those are particularly used which have a high content of water of crystallization. Aluminum sulfate hydrate, in particular the compounds $Al_2(SO_4)_3 \cdot 16H_2O$ and $Al_2(SO_4)_3 \cdot 18H_2O$ having the particularly high water of crystallization content of 16 and 18 moles of H_2O /mole of $Al_2(SO_4)_3$, respectively, is particularly preferred.

15 A further variant for the preparation of aluminoxane is to dissolve trialkylaluminum, preferably trimethylaluminum, in the suspending agent initially introduced into the polymerization reactor and then to react the aluminum compound with water.

20 Besides the above-outlined processes for the preparation of aluminoxanes, there are others which can be used.

25 Irrespective of the manner of preparation, all aluminoxane solutions have in common a varying content of unreacted trialkylaluminum, which is in free form or in the form of an adduct. This content has an effect, as yet not explained accurately, on the catalytic effectiveness, which varies depending on the metallocene compound employed.

30 It is possible to pre-activate the metallocene using an aluminoxane of the formula II and/or III before using the polymerization reaction. This considerably increases the polymerization activity and improves the grain morphology.

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5 The preactivation of the transition-metal compound is carried out in solution. It is preferred here to dissolve the metallocene in a solution of the aluminoxane in an inert hydrocarbon. Suitable inert hydrocarbons are aliphatic or aromatic hydrocarbons. Toluene is preferably used.

10 The concentration of the aluminoxane in the solution is in the range from about 1% by weight to the saturation limit, preferably from 5 to 30% by weight, in each case relative to the total solution. The metallocene can be employed in the same concentration, but it is preferably employed in an amount of from 10^{-4} - 1 mole per mole of aluminoxane. The pre-activation time is 5 minutes to 60 hours, preferably 5 to 60 minutes. The pre-activation is carried out at a temperature of from -78°C to 100°C, preferably 0 to 70°C.

15 A significantly longer preactivation is possible, but normally has neither an activity-increasing nor activity-reducing effect, but may be entirely appropriate for storage purposes.

20 The polymerization is carried out in a known manner in solution, in suspension or in the gas phase, continuously or batchwise, in one or more steps, at a temperature of from -60 to 300°C, preferably -30 to 120°C, in particular 50 to 90°C.

25 The overall pressure in the polymerization system is 0.5 to 200 bar. The polymerization is preferably carried out in the industrially particularly interesting pressure range of from 5 to 60 bar. The metallocene compound is used here in a concentration, relative to the transition metal, of from 10^{-3} to 10^{-1} , preferably 10^{-4} to 10^{-2} mole of transition metal per dm³ of solvent or per dm³ of reactor volume. The aluminoxane is used in a concentration of from 10^{-3} to 10^{-1} mole, preferably 10^{-4} to 10^{-2} mole per dm³ of solvent or per dm³ of reactor volume. IN

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principle, however, higher concentrations are also possible.

5 If the polymerisation is carried out as a suspension or solution polymerisation, a solvent which is inert towards Siegler catalysts is used, i.e. an aliphatic or aromatic hydrocarbon. Aliphatic hydrocarbons are preferred, such as, for example, butane, pentane, hexane, heptane, isooctane, cyclohexane, methylcyclohexane or petroleum or hydrogenated diesel oil fractions.

10 Besides homopolymerization of ethylene, the catalyst systems according to the invention are employed for the copolymerization of ethylene with a 1-olefin having 3 to 20 carbon atoms. Examples of such 1-olefins are propylene, 1-butene, 1-hexene, 4-methyl-1-pentene and 1-octene.

15 The molecular weight of the polymer can be regulated in a known manner, preferably using hydrogen.

20 The polymerisation can have any desired duration, since the catalyst system to be used according to the invention only exhibits a slight time-dependent decrease in polymerization activity.

25 Depending on the structure of the complex, the use of these complexes gives, with high activities, polyethylene and ethylene-1-olefin copolymers having a narrow molecular-weight distribution (polydispersity) in a broad molecular-weight range, in particular products having high molecular weights which are suitable for processing by injection molding and extrusion and in particular have high stretching capacity of the polymer melt, and permits the production of very varied grain morphologies of the product, such as high and low bulk densities, extremely small and extremely large mean grain diameters and various grain shapes. The variety of grain morphologies which can be achieved opens up various possibilities for the use of such polyethylene powders in sintering processes.

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The examples below are intended to illustrate the invention.

The abbreviations have the following meanings:

5 η_{sp} = viscosity number in cm²/g,
 M_w = weight average molecular weight in g/mol,
 M_w/M_n = molecular weight distribution determined by gel
permeation chromatography (GPC),
d₅₀ = mean grain diameter in μ m.

10 The pressure data in the examples are in bar of super-
atmospheric pressure.

The densities of the copolymers have been determined in
accordance with DIN 53479, method A.

All the operations below were carried out under a protective
gas using absolute solvents.

15 Example 1:

Preparation of dimethylsilylbis(1-indenyl)

20 80 cm³ (0.20 mol) of a 2.5 molar solution of n-butyl-
lithium in hexane were added with ice cooling to a solu-
tion of 30 g (0.23 mol) of indene (technical grade, -91%)
which has been filtered through aluminum oxide. The batch
25 was stirred for a further 15 minutes at room temperature,
and the orange solution was added via a hollow needle
over the course of 2 hours to a solution of 13.0 g
(0.10 mol) of dimethyldichlorosilane (99%) in 30 cm³ of
diethyl ether. The orange suspension was stirred over-
night and washed by shaking three times with 100-150 cm³
30 of water. The yellow organic phase was dried twice over
sodium sulfate and evaporated in the reaction evaporator.
The orange oil which remained was kept at 40°C for 4 to
5 hours in an oil-pump vacuum and freed from excess
indene, during which a white precipitate deposited.
Addition of 40 cm³ of methanol and crystallization at
-35°C gave a total of 25.4 g (71%) of the compound

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dimethylsilyl bis(1-indenyl) as a white to beige powder (2 diastereomers). b.p. 79-81°C.

Example 2:

Preparation of rac-dimethylsilyl bis(1-indenyl) siliconium dichloride (metallocene A)

15.5 cm³ (38.7 mmol) of a 2.5 molar hexane solution of butyllithium were added slowly at room temperature to a solution of 5.6 g (19.4 mmol) of dimethylsilyl bis(1-indenyl) in 40 cm³ of THF. 1 hour after completion of the addition, the deep red solution was added dropwise over the course of 4-5 hours to a suspension of 7.3 g (18.4 mmol) of $\text{SiCl}_4\text{-THF}$ in 60 cm³ of THF. After the mixture had been stirred for 3 hours, the orange precipitate was filtered off with suction via a glass frit and recrystallized from CH_2Cl_2 , to give 1.0 g (11%) of the metallocene A in the form of orange crystals, which gradually decompose from 200°C.

Correct elementary analyses. The EI mass spectrum exhibited $M^+ \sim 448$. ^1H NMR spectrum (CDCl_3): 7.04-7.50 (m, 8, aron. H), 6.90 (dd, 2, β -Ind H), 6.08 (d, 2, α -Ind H), 1.12 (s, 6, SiCH_3).

Example 3:

Preparation of rac-diphenylsilyl bis(1-indenyl) siliconium dichloride (metallocene B)

A solution of 20 g (49.5 mmol) of $(\text{C}_6\text{H}_5)_2\text{Si}(\text{Ind})_2$, prepared from $(\text{C}_6\text{H}_5)_2\text{SiCl}$, and indenyllithium analogously to Example 1, in 200 cm³ of diethyl ether was reacted at 0°C with 40 cm³ (100 mmol) of butyllithium (2.5 molar in hexane). After the mixture had been stirred at room temperature for 2 hours, the solvent was stripped off, and the residue was stirred with 100 cm³ of hexane and filtered off. The dilithio salt was dried in an oil-pump vacuum and added to a suspension of 11.3 g (43.5 mmol) of SiCl_4 in 130 cm³ of CH_2Cl_2 at -78°C. The mixture was stirred

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overnight and allowed to warm to room temperature. The 5
rod solution was concentrated, and the precipitate which
deposited was filtered off via a frit. Extraction with
toluene gave 2.0 g (74%) of metallocene E as an orange
powder. Correct elementary analyses. ^1H NMR spectrum
(CDCl₃): 6.8-8.2 (m, 18, arom. H), 7.03 (dd, 2, β -Ind H),
5.30 (d, 2, α -Ind H).

Example 4:

Preparation of rac-dimethylsilyl bis(1-(3-methylindenyl))-
10
zirconium dichloride (metallocene G)

A solution of 4.85 g (15.5 mmol) of (CH₃)₂Si(MeInd)₂, prepared 15
analogously to Example 1 from (CH₃)₂SiCl₂ and 3-
methylindenyllithium, was reacted with 12.5 cm³
(30.5 mmol) of butyllithium and 5.84 g (15.5 mmol) of
20
EtCl₂.²⁷ Analogously to Example 2. After the solvent had
been stripped off, the residue was extracted with toluene.
The precipitate which deposited from toluene on
concentration and cooling was recrystallized from CHCl₃,
to give 800 mg (10%) of metallocene G in the form of
orange-red crystals. Correct elementary analyses. ^1H NMR
Spectrum (CDCl₃): 7.0-7.5 (m, 8, arom. H), 5.21 (s, 2,
 α -Ind H), 2.30 (s, 6, Ind CH₃), 1.07 (s, 6, SiCH₃).

Example 5:

Preparation of dimethylsilyl bis(1-(3-trimethylsilyl)-
25
cyclopentadienyl)zirconium dichloride (metallocene F)

A solution of 3.9 g (11.7 mmol) of (CH₃)₂Si(CH₃)₂Si(Cp)₂, prepared 30
from Li₂[(CH₃)₂Si(Cp)₂] and (CH₃)₂SiCl₂, was reacted
analogously to Example 3 with 9.4 cm³ (23.4 mmol) of butyllithium
and 4.4 g (11.7 mmol) of EtCl₂.²⁷ After the solvent had been stripped off, the residue was extracted
with diethyl ether. The residue remaining after the 35
diethyl ether had been stripped off was recrystallized from CHCl₃, to give 0.8 g (14%) of the complex as beige
crystals. Correct elementary analyses. ^1H NMR spectrum
(CDCl₃): 6.93 (dd, 2, CpH), 6.12 (t, 2, CpH), 5.86 (t, 2,

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CpM), 0.72 (s, 6, Si(C₂H₅)₂), 0.25 (s, 9, Si(CH₃)₃). The NMR spectrum showed that metallocene V was in the form of an isomer mixture (43% of the rac-isomer, 57% of the meso-isomer).

5 Example 6:

Preparation of isopropyl(1-indanyl)cyclopentadienylaluminum dichloride (metallocene F)

10 19 cm³ (47.3 mmol) of butyllithium (2.5 molar in hexane) were added at room temperature to a solution of 6.0 g (47 mmol) of indane (91%) in 100 cm³ of diethyl ether. After 1 hour, this solution was added to a solution of 6,6-dimethylfulvalene in 100 cm³ of diethyl ether at -78°C. After the mixture had been stirred at room temperature for 16 hours, the orange solution was diluted with 400 cm³ of diethyl ether, and 100 cm³ of water were added. The yellow organic phase was then washed twice with water, dried over Na₂SO₄, and evaporated. The brown oil remaining was chromatographed on 400 g of silica gel 60. Using benzene + 7% of methylene chloride, a total of 7.2 g (68%) of the compound isopropyl(1-indanyl)cyclopentadienyl were eluted as a yellow oil (2 isomers). 20 cm³ (70 mmol) of butyllithium (2.5 molar in hexane) were added at 0°C to a solution of 7.1 g (32 mmol) of this compound in 100 cm³ of diethyl ether. After the mixture had been stirred at room temperature for 2 hours, the yellow precipitate was filtered off via a glass frit and washed with hexane/diethyl ether (1:1). After the precipitate had been dried in an oil-pump vacuum the pale yellow powder was added at -78°C to a suspension of 7.5 g (32 mmol) of ZnCl₂ in 100 cm³ of methylene chloride. The mixture was slowly warmed to room temperature, stirred at room temperature for 30 minutes and filtered via a glass frit, and the solid was washed several times with methylene chloride. The yellow filtrate was concentrated until crystallization occurred. At -35°C, a total of 2.4 g (19%) of the complex rac-[(CH₃)₂C(Ind)Cp]ZnCl₂ crystallized in the form of yellow-orange crystals. Correct elementary analyses.

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¹H NMR spectrum (CDCl₃): 6.90-7.75 (m, 4, arom. H), 6.05 (dd, 1, β -Ind-H), 8.12 (m, 3, Cp-H), 6.12 (d, 1, α -Ind-H), 5.82, 5.70 (2 x q, 2 x 1, Cp-H), 2.20; 1.95 (2 x s, 2 x 3, CH₃).

5 Example 7a

Preparation of diphenylmethylene(β -fluoromethyl)cyclopentadienylaluminum dichloride (metallocene Q)

10 12.5 cm³ (30.7 mmol) of a 2.3 molar hexane solution of n-butyllithium were slowly added at room temperature to a solution of 5.10 g (30.7 mmol) of fluorene in 50 cm³ of THF. After 40 minutes, 7.07 g (30.7 mmol) of diphenylfulvene were added to the orange solution, and the mixture was stirred overnight. 50 cm³ of water were added to the dark red solution, whereupon the solution became yellow, and the solution was extracted using ether. The ether phase, dried over MgSO₄, was concentrated and left to crystallize at -95°C, to give 5.1 g (42%) of 1,1-cyclopentadienyl(β -fluoromethyl)diphenylmethane as a beige powder.

20 2.0 g (5.0 mmol) of the compound were dissolved in 20 cm³ of THF, and 6.4 cm³ (10 mmol) of a 1.6 molar solution of butyllithium in hexane were added at 0°C. After the mixture had been stirred at room temperature for 15 minutes, the solvent was stripped off, and the red residue was dried in an oil-pump vacuum and washed several times with hexane. After the residue had been dried in an oil-pump vacuum, the red powder was added at -76°C to a suspension of 1.16 g (5.0 mmol) of SrCl₂. The batch was slowly warmed and stirred for a further 3 hours at room temperature. The pink suspension was filtered via a G3 frit. The pink residue was washed with 20 cm³ of CH₂Cl₂, dried in an oil-pump vacuum and extracted with 120 cm³ of toluene. Stripping off of the solvent and drying in an oil-pump vacuum gave 0.55 g of the aluminon complex (metallocene Q) in the form of a pink crystal powder.

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The orange-red filtrate of the reaction batch was evaporated and left to crystallize at -35°C. A further 0.34 g of the complex crystallizes from CH_2Cl_2 . Total yield 1.0 g (36%). Correct elementary analyses. The mass spectrum exhibited $M^+ = 556$. ^1H NMR spectrum (100 MHz, CDCl_3): 6.90-8.25 (m, 16, Flu-H, Ph-H), 6.40 (m, 2, Ph-H), 5.37 (t, 2, Cp-H), 5.80 (t, 2, Cp-H).

Example 8-10:

The metallocenes C, D, E, F, H, I, L, M, N and R of Table 2 were prepared analogously to Examples 1 and 2. The dimethylchlorosilane in Example 1 was replaced here by appropriate ether dihalogen compounds, which are shown in Table 1. In the case of complexes substituted on the 5-membered ring (metallocenes N and R), an indene correspondingly substituted on the 5-membered ring was employed (analogously to Example 4). In the case of the hafnium complex metallocene R, HfCl_4 in Example 2 was replaced by MgCl_2 .

Table 1

Example	Metallocene	Dihalogen compound
8	C	phenylmethyldichlorosilane
9	D	phenylvinyldichlorosilane
10	E	dimethyl dichlorogermylum
11	F	cycloheximethylenedichlorosilane
12	H	1,1,2,2-tetramethyl-1,2-dichlorodisilane
13	I	1,2-bis(chlorodimethylsilyl)-ethane
14	L	1,2-dibromoethane
15	M	1,3-dibromopropane
16	M	1,2-dibromoethane
17	O	1,2-dibromosthene
18	R	1,1,2,2-tetramethyl-1,2-dichlorosilane

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Example 19:

Preparation of phenylmethylmethylcyclo(9-fluorenyl)cyclopentadienylhafnium dichloride (metallocene 2)
The metallocene 2 was prepared analogously to Example 7.
5 However, the diphenylfulvene and HfCl_4 in Example 7 were replaced by phenylmethylfulvone and HfCl_4 .

Example 20:

10 10 dm^3 of a petroleum ether (boiling range 100-120°C) were introduced at 20°C into a dry 16 dm^3 reactor which had been flushed with nitrogen. The gas space of the reactor was then flushed free from nitrogen by injecting 2 bar of ethylene and releasing the ethylene, and repeating this operation 4 times. 1 bar of hydrogen was then injected, and 30 cm^3 of a toluene solution of methylaluminoxane 10.5% by weight of methylaluminoxane, molecular weight 750 g/mol according to cryoscopic determination). The reactor contents were heated to 60°C over the course of 15 minutes with stirring. The overall pressure was then increased to 7 bar by introducing ethylene with stirring at 250 rpm. At the same time, 3.1 mg of metallocene 2 were dissolved in 20 cm^3 of a toluene solution of methylaluminoxane (concentration and quality as above) and pre-activated by being left to stand for 15 minutes. The solution was then introduced into the reactor. The polymerization system was warmed to a temperature of 65°C and then kept at this temperature for 1 hour by means of suitable cooling. The overall pressure was kept at 7 bar during this time by corresponding supply of ethylene. 25 160 g of polyethylene were obtained.

30 The following values were determined on the products:
VM = 132 cm^3/g , bulk density: 320 g/ dm^3

Examples 21-42:

35 The procedure was in each case analogous to Example 20, but the following parameters were varied:
- metallocene type